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	Filing Date		2006-08-30		
	First Named Inventor	Philip	nilip Wilson		
	Art Unit				
	Examiner Name	Unkn	known		
	Attorney Docket Numb	er	065435-9083 US00		

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1999-09-16

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1 6613787

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	1	ANDERSEN, H.S. et al., "Discovery and SAR of a novel selective and orally bloavailable nonpeptide classical competitive inhibitor class of protein-tyrosine phosphatase 18," J. Med. Chem. (2002) 45:4443-4459	
	2	BERGE, S.M. et al., "Pharmacousical Salts," J. Pharm. Sci. (1977) 66(1):1-19	
	3	BRIEFN, C.A. et al., "Alternative heterocycles for DNA recognition: the benzimidazole/imidazole/pair," Chem. Eur. J. (2003) 9-2110-2122	
	4	DANNLEY, R.L. et al., "Free radical aromatic substitution. IV. The reaction of acyl peroxides with berzotrihalides," J. Am. Chem. Soc. (1954) 78:4543-4546	
	5	GREENE, T.W. et al., Protective Groups in Organic Synthesis, 2nd Edition, Wiley Publishing (1991) Chapter 7, 315-345	
	6	GREGSON, S.J. et al., "Synthesis of the first examples of A-CB/C-C2 amide-linked pyrrolo[2,1-c][1,4]benzodzepine dimers," Blorg. Med. Chem. Lett. (2009) 13:2277-2280	
	7	NISHIWAVI, E. et at, "Efficient synthesis of oligo-n-methylpymolecurboxamides and related compounds," Heterocycles (1989) 27(8): 1945-1992	
	8	PERJESSY, A. et al., "Application of the Seth-Paul-Van Duyse equation - III. Transmission of polar effects by the furan ring," Tetrahedron (1975) 31:2936-2939	
	9	QIAN, Y. et al., "Design and synthesis of non-peptide Ras CAAX mimetics as potent famesyltransferase inhibitors," J. Med. Chem. (1988) 39:217-223	
	10	QUAN, M.L. et al., "Blaryl substituted alkythoronate esters as thrombin inhibitors," Blorg. Med. Chem. Lett. (1997) 7 (13):1595-1800	

11	RENNEBERG, D. et al., "Imidazopyridinelpyrrole and hydroxybenzimidazole/pyrrole pairs for DNA minor groove recognition." J. Am. Chem. Soc. (2003) 126:5707-5716	
12	WOODS, C.R. et al., "Synthesis and DNA binding properties of immodiscetic acid-linked polyamides: characterization of cooperative extended 2:1 side-by-side parallel binding," J. Am. Chem. Soc. (2002) 124:10878-10882	

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